

Sub D2

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8. A vector according to ~~any one of the preceding claims~~, wherein the one or more cleavage site(s) is/are located at a site(s) on the vector which avoids steric hindrance of binding by said restriction enzyme or functional portion thereof.

9. A vector according to ~~any one of the preceding claims~~, further comprising a third nucleotide sequence encoding a ribozyme targetted against mRNA produced from the said second nucleotide sequence encoding the restriction enzyme or functional portion thereof.

10. A vector according to ~~any one of the preceding claims~~, wherein the second promoter is selected from the group consisting of the *placZ* promoter, the *placUV5* promoter and the T7 RNA polymerase promoter.

11. A vector according to claim 10, wherein the second promoter is the T7 RNA polymerase promoter.

12. A vector according to claim 11, further comprising an additional nucleotide sequence encoding T7 RNA polymerase operably linked to a third promoter sequence, said third promoter sequence being inducible.

13. A host cell transformed with a suicide expression vector according to ~~any one of the preceding claims~~.

14. A host cell according to claim 13, wherein said host cell is a bacterium or yeast.

15. A method of expressing a heterologous peptide, polypeptide or protein in a selected host cell, comprising;

(i) transforming said host cell with a suicide expression vector according to ~~any one of claims 1 to 12~~;

(ii) culturing said transformed host cell under suitable conditions for the expression of the said heterologous peptide, polypeptide or protein, and

(iii) thereafter inducing expression of the restriction enzyme or functional portion thereof to bring about cleavage of the said suicide expression vector.

16. A method according to claim 15, wherein the host cell is a bacterium or yeast.

sub D6 > 5 17. A method for the production of a microorganism vector which contains recombinant peptide, polypeptide or protein but no recombinant DNA, comprising;

9 (i) transforming said microorganism with a suicide expression vector according to any one of claims 1 to 12;

10 (ii) culturing said transformed microorganism under suitable conditions for the expression of the said heterologous peptide, polypeptide or protein, and (iii) thereafter inducing expression of the restriction enzyme or functional portion thereof to bring about cleavage of the said suicide expression vector.

15 18. A method according to claim 17, wherein the microorganism is a bacterium or yeast.

19. A microorganism vector produced by the method according to claim 17 or 18;

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